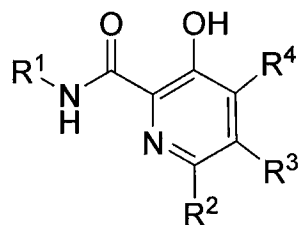


Listing of Claims

The listing of claims will replace all prior versions, and listings, of claims in the application:

1. (Currently Amended) A compound of the Formula:



wherein:

R¹ is selected from:

- a) hydrogen,
- b) aryl, heterocycle, C₃-C₁₀ cycloalkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, and
- c) C₁-C₆ alkyl, unsubstituted or substituted with 1 to 5 substituents selected from:
 - 1) aryl, unsubstituted or substituted with 1 to 5 substituents selected from:
 - i) C₁-C₆ alkyl, unsubstituted or substituted with 1-3 fluoro,
 - ii) C₃-C₆ cycloalkyl,
 - iii) C₂-C₆ alkynyl,
 - iv) OR¹⁰,
 - v) aryl,
 - vi) heterocycle,
 - vii) CN, and
 - viii) halo;
 - 2) heterocycle, unsubstituted or substituted with 1 to 5 substituents selected from:
 - i) C₁-C₆ alkyl, unsubstituted or substituted with 1-3 fluoro,
 - ii) -OR¹⁰,

- iii) aryl, and
- iv) halo;
- 3) C₃-C₁₀ cycloalkyl,
- 4) C₂-C₆ alkenyl,
- 5) C₂-C₆ alkynyl,
- 6) -OR¹⁰,
- 7) -S(O)_mR¹¹,
- 8) -NR⁶-C(O)R⁷,
- 9) -C(O)-N(R⁶)(R⁷),
- 10) -CN,
- 11) -NR⁶-C(O)-N(R⁶)(R⁷),
- 12) -C(O)-OR¹⁰,
- 13) halo, and
- 14) -N(R⁶)(R⁷);

R² is selected from:

- a) -NR⁶-C(O)R⁷,
- b) -NR⁶-S(O)₂R⁷, and
- c) -NR⁶-S(O)₂-N(R⁶)(R⁷);

R³ and R⁴ are independently selected from:

hydrogen, aryl, heterocycle, halo, C₁-C₆ alkyl, C₃-C₁₀ cycloalkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, C₁-C₄ haloalkyl, R¹⁰O-, R¹¹S(O)_m-, R⁶C(O)-NR⁷-, CN, (R⁶)(R⁷)N-C(O)-(NR⁶)-, (R⁶)(R⁷)-N-C(O)-, R¹⁰C(O)-, R¹⁰OC(O)-, and N(R⁶)(R⁷); or

wherein R³ and R⁴ are optionally joined to form a saturated or unsaturated ring, containing 0-3 heteroatoms, wherein said ring is phenyl, ~~pyridyl~~, pyrimidinyl, pyrazinyl, thiophenyl, furanyl, imidazolyl, thiazolyl, oxazolyl, and triazolyl, ~~as well as~~ and partially saturated analogues thereof, said ring optionally substituted with one or more of:

aryl, heterocycle, C₁-C₆ alkyl, C₃-C₁₀ cycloalkyl, C₂-C₆ alkynyl, R¹⁰O-, R¹¹S(O)_m-, R⁶C(O)N R⁷-, -R⁶S(O)₂NR⁷-, (R⁶)(R⁷)N-C(O)-, CN, R¹⁰OC(O)-, F, and -N(R⁶)(R⁷);

R⁶ and R⁷ are independently selected from hydrogen, C₁-C₆ alkyl, C₃-C₁₀ cycloalkyl, heterocycle, aryl, unsubstituted or substituted ~~with one~~ with one or more of:

- a) C₁-C₄ alkyl,
- b) C₁-C₄ alkoxy,
- c) aryl or heterocycle,
- d) halo,
- e) -OR¹⁰, and
- f) -N(R¹⁰)₂;

wherein R⁶ and R⁷ may be joined to form a ring;

R¹⁰ is independently selected from hydrogen, C₁-C₆ alkyl, -CF₃, C₃-C₁₀ cycloalkyl, benzyl, and aryl;

R¹¹ is independently selected from C₁-C₆ alkyl, and aryl;

m is 0, 1, or 2;

or a ~~and~~ pharmaceutically acceptable salt ~~salts~~ and individual diastereomers thereof.

2. (Original) The compound according to Claim 1, wherein R¹ is -CH₂-aryl, unsubstituted or substituted with 1-3 substituents selected from: fluoro, chloro, bromo, iodo and methyl.

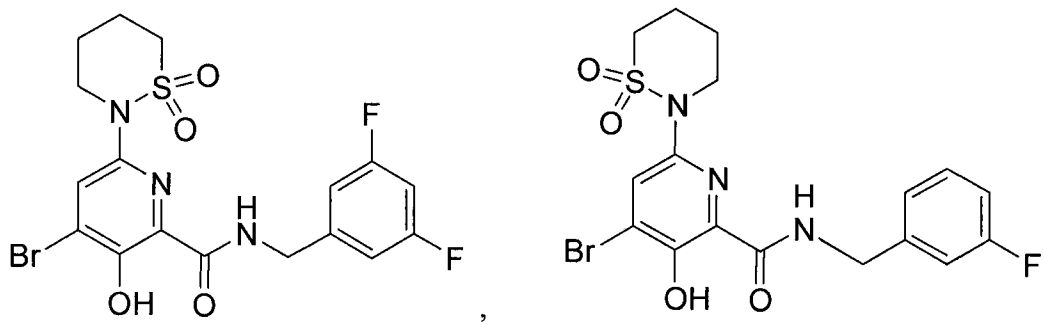
3. (Original) The compound according to Claim 1, wherein R¹ is benzyl, substituted with 1-3 fluoro.

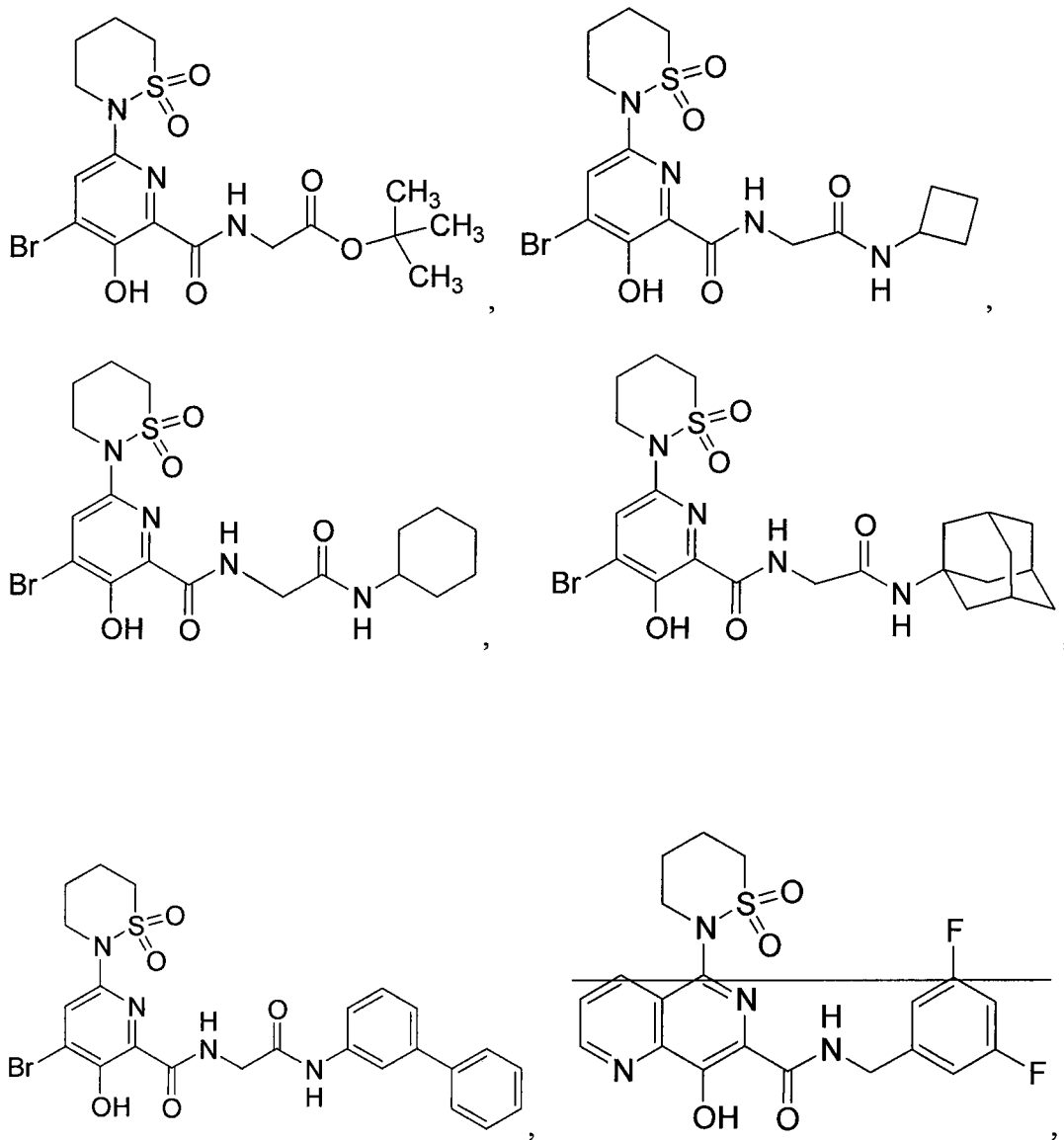
4. (Original) The compound according to Claim 1, wherein R¹ is -CH₂C(O)OR¹⁰.

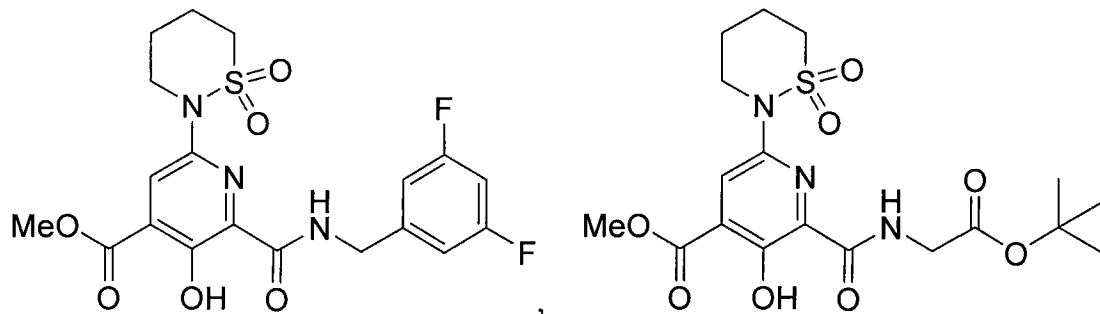
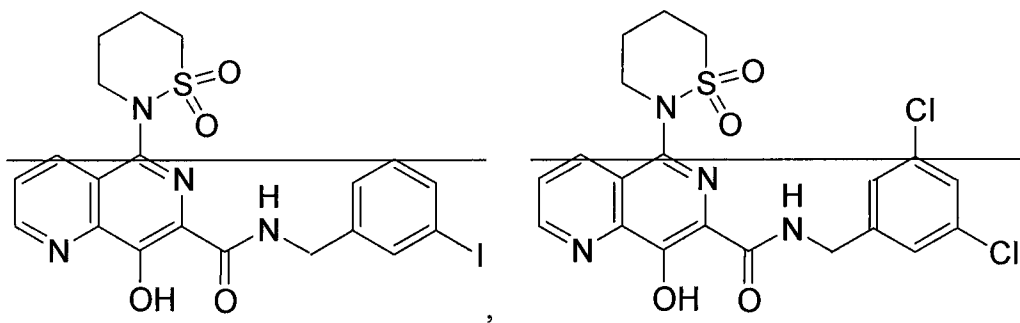
5. (Original) The compound according to Claim 1, wherein R¹ is -CH₂C(O)OC(CH₃)₃.

6. (Original) The compound according to Claim 1, wherein R¹ is -CH₂C(O)NHR⁶.

7. (Original) The compound according to Claim 1, wherein R¹ is -CH₂C(O)NH(C₄-C₁₀ cycloalkyl).
8. (Original) The compound according to Claim 1, wherein R¹ is -CH₂C(O)NH-aryl.
9. (Original) The compound according to Claim 1, wherein R² is -NR⁶-S(O)₂R⁷.
10. (Original) The compound according to Claim 1, wherein R³ is hydrogen.
11. (Currently Amended) The compound according to Claim 1, wherein R³ and R⁴ are joined to form a ring selected from: phenyl, ~~pyridyl~~, pyrimidinyl and pyrazinyl.
12. (Canceled)
13. (Original) The compound according to Claim 1, wherein R⁴ is bromo.
14. (Original) The compound according to Claim 1, wherein R⁴ is -C(O)OR¹⁰.
15. (Currently Amended) A compound selected from:







and pharmaceutically acceptable salts and individual diastereomers thereof.

16. (Currently Amended) A pharmaceutical composition which comprises a pharmaceutically acceptable ~~an~~ inert carrier and the compound of Claim 1.

17. (Canceled)

18. (Previously Amended) A method for treating, controlling, ameliorating or reducing the risk of headache in a mammalian patient in need of such which comprises administering to the patient a therapeutically effective amount of the compound of Claim 1.

19 24. (Canceled)

25. (Previously Presented) The method of claim 18, wherein the headache is migraine headache or cluster headache.